WHAT IS CLAIMED IS:

1	1. A quick disintegrating tablet in buccal cavity, said quick disintegrating
2	tablet comprising:
3	a) a plurality of drug-containing particles, wherein each particle comprises a
4	bitter tasting drug and/or a drug of inferior fluidity and a pharmaceutical preparation carrier,
5	wherein each particle has a mean diameter of approximately 50 to approximately 250 μm and
6	an apparent specific gravity of approximately 0.5 to approximately 1.2; and
7	b) a saccharide.
1	2. The quick disintegrating tablet in buccal cavity of claim 1, wherein the
2	drug of inferior fluidity has an angle of repose of 41° ~ 90°.
1	The quick disintegrating tablet in buccal cavity of claim 1, wherein the
2	pharmaceutical preparation carrier is 1 or 2 or more selected from the group consisting of
3	water-insoluble polymers, gastrosoluble polymers, enterosoluble polymers, wax-like
4	substances and saccharides.
1	4. The quick disintegrating tablet in buccal cavity of claim 3, wherein the
1	pharmaceutical preparation carrier is a water-insoluble polymer.
2	pharmaceutical preparation carrier is a water-insoluble polymer.
1	5. The quick disintegrating tablet in buccal cavity of claim 4, wherein the
2	water-insoluble polymer is a water-insoluble cellulose ether or a water-insoluble acrylic acid
3	copolymer.
1	6. The quick disintegrating tablet in buccal cavity of claim 1, wherein the
2	amount of pharmaceutical preparation carrier added is about 0.05 to about 3 parts by weight
3	per 1 part by weight bitter tasting drug and/or drug of inferior fluidity.
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1	7. The quick disintegrating tablet in buccal cavity of claim 1, wherein the
2	saccharide is a granulation product obtained by spraying to coat and/or granulate a saccharide
3	of low moldability using a saccharide of high moldability as a binder.
1	8. The quick disintegrating tablet in buccal cavity of claim 7, wherein the
2	saccharide of low moldability is 1 or 2 or more selected from the group consisting of lactose,
3	mannitol, glucose, sucrose, xylitol, and erythritol.

9. The quick disintegrating tablet in buccal cavity of claim 7, wherein the saccharide of high moldability is 1 or 2 or more selected from the group consisting of maltose, maltitol, sorbitol, trehalose, and lactosucrose.

- The quick disintegrating tablet in buccal cavity of claim 1, wherein the
 mean particle diameter of the plurality of drug-containing particles is approximately 50 μm to
 approximately 150 μm.
- 1 11. The quick disintegrating tablet in buccal cavity of claim 1, wherein the apparent specific gravity of the plurality of drug-containing particles is approximately 0.5 ~ approximately 1.
- 1 12. A drug-containing particle, wherein said drug containing particle has a
 2 mean particle diameter of approximately 50 to approximately 250 μm and an apparent
 3 specific gravity of approximately 0.5 to approximately 1.2, and comprises a bitter tasting
 4 drug and a water-insoluble polymer.
- 1 13. A drug-containing particle, wherein said drug containing particle has a mean particle diameter of approximately 50 to approximately 250 μm and an apparent specific gravity of approximately 0.5 to approximately 1.2, and comprises a drug of inferior fluidity and a saccharide.
 - 14. A method for manufacturing a quick disintegrating tablet in buccal cavity, said quick disintegrating tablet comprising a drug and a saccharide, said method comprising the steps of:
 - (a) dissolving a bitter tasting drug and/or a drug of inferior fluidity and a pharmaceutical preparation carrier to form a mixture that is dissolved and suspended to approximately 30 to approximately 70 w/w% in terms of solid concentration in a solvent that is pharmaceutically acceptable to prepare a suspension for spray drying;
 - (b) spray drying said suspension using a rotating disk-type spray dryer, with the disk rotating at a speed of approximately 5,000 to approximately 15,000 rpm to prepare the drug-containing particles; and
- 11 (c) mixing the drug-containing particles with a saccharide to form a mixture that is molded.

- 1 15. The method for manufacturing a quick disintegrating tablet in buccal 2 cavity of claim 14, wherein said saccharide is a granulation product obtained by spraying to 3 coat and/or granulate a saccharide of low moldability using a saccharide of high moldability 4 as a binder.
- 1 16. A method for manufacturing a quick disintegrating tablet in buccal 2 cavity of claim 14, wherein (d) the process of moistening and drying is further performed in 3 succession to process (c) on the molding obtained under at least the pressure needed to retain 4 tablet form.
- 1 17. The method for manufacturing a quick disintegrating tablet in buccal 2 cavity of claim 14, wherein the solid concentration in step (a) is approximately 40 to 3 approximately 70 w/w%.
- 1 18. The method for manufacturing a quick disintegrating tablet in buccal 2 cavity of claim 14, wherein the rotating speed of the rotating disk in process (b) is 3 approximately 6,000 to approximately 12,000 rpm.
- 1 19. The method for manufacturing a quick disintegrating tablet in buccal cavity of claim 14, wherein a bitter tasting drug and/or a drug of inferior fluidity whose particle diameter has been brought to approximately 5 to approximately 100 µm is used in process (a).

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20. 1 A quick disintegrating tablet in buccal cavity, which is manufactured 2 by the method of claim 14.